

1621  
JFW



PATENT

THE UNITED STATES PATENT AND TRADEMARK OFFICE  
(Case No. 99,139-G)

|  |   |                      |
|--|---|----------------------|
| In the Application of:                       | ) |                      |
|  | ) |                      |
| Kolb et al.                                  | ) |                      |
|  | ) | Examiner: P. Zucker  |
| Serial No.: 10/797,458                       | ) |                      |
|  | ) | Group Art Unit: 1621 |
| Filing Date: March 10, 2004                  | ) |                      |
|  | ) |                      |
| For: Novel PPAR- $\gamma$ Agonists as Agents | ) |                      |
| for the Treatment of Type II                 | ) |                      |
| Diabetes                                     | ) |                      |

Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

TRANSMITTAL LETTER

In regard to the above-identified patent application:

1. We are transmitting herewith the attached:
  - a. Response to the Office Action mailed November 2, 2004
  - b. Postcard
2. Please charge any additional fees or credit over-payments to the Deposit Account No.13-2490.
4. **CERTIFICATE OF MAILING UNDER 37 CFR § 1.8:** The undersigned hereby certifies that this Transmittal Letter and the paper(s), as described in paragraph 1 hereinabove, are being deposited with the United States Postal Service with sufficient postage as First Class Mail in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA on this 3rd day of January, 2005.

Dated: January 3, 2005

By: Stephen H. Docter  
Stephen H. Docter  
Reg. No. 44,659



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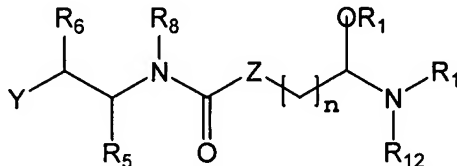
**RESPONSE TO THE OFFICE ACTION MAILED NOVEMBER 2, 2004**

Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

Dear Sir:

Responsive to the Office Action Mailed November 2, 2004 in the above-mentioned case, Applicants can not elect a species as set forth in present claim 3, as all compounds therein are the subject matter of issued claims found in the parent case, U.S. Patent No. 6,713,514, issued from U.S.S.N. 09/552,477. As a result, Applicants herewith propose and election of invention within the subject matter of claim 1 as follows.

A compound of the formula



or the pharmaceutically acceptable non-toxic salts thereof wherein:

Z is a 5 or 6 membered aryl or heteroaryl ring optionally substituted with up to three groups selected from lower alkyl, halogen or lower alkoxy;

n is 1 or 2;

R<sub>1</sub> and R<sub>12</sub> are the same or different and represent hydrogen, lower alkyl, SO<sub>2</sub>(R<sub>10</sub>), or cycloalkyl optionally substituted with one, two, three or four groups independently selected from halogen, trifluoromethyl, trifluoromethoxy, cyano, nitro, carboxyl, alkoxy, alkoxy, alkyl, carboxyl, hydroxy, lower alkyl, lower alkoxy, amino, or mono or dialkylamino where each alkyl portion is lower alkyl, or

aryl, heteroaryl, arylalkyl, or heteroarylalkyl, where the ring portion of each is optionally substituted with one, two, three or four groups independently selected from halogen, trifluoromethyl, trifluoromethoxy, cyano, nitro, carboxyl, alkoxy, alkoxy, alkyl, carboxyl, hydroxy, lower alkyl, lower alkoxy, amino, or mono or dialkylamino where each alkyl portion is lower alkyl;

R<sub>10</sub> is hydrogen or lower alkyl, or aryl, heteroaryl, arylalkyl or heteroarylalkyl, where the ring portion of each is optionally substituted with one, two or three groups independently selected from halogen, trifluoromethyl, trifluoromethoxy, cyano, nitro, carboxyl, alkoxy, alkoxy, alkyl, carboxyl, hydroxy, lower alkyl, lower alkoxy, amino, or mono or dialkylamino where each alkyl portion is lower alkyl;

R<sub>9</sub> is H or lower alkyl;

Y is hydrogen, NR<sub>1</sub>R<sub>12</sub>, OR<sub>1</sub>, CH<sub>2</sub>R<sub>1</sub>, SR<sub>1</sub>, SOR<sub>1</sub> or SO<sub>2</sub>R<sub>1</sub>; and

R<sub>5</sub>, R<sub>6</sub> and R<sub>8</sub>, are the same or different and represent hydrogen, lower alkyl, R<sub>10</sub>C=O,

R<sub>10</sub>SO<sub>2</sub>, or

cycloalkyl optionally substituted with one, two, three or four groups

independently selected from halogen, trifluoromethyl, trifluoromethoxy,

cyano, nitro, carboxyl, alkoxycarboxy, alkylcarboxy, hydroxy, lower

alkyl, lower alkoxy, amino, or mono or dialkylamino where each alkyl

portion is lower alkyl, or

aryl, heteroaryl, arylalkyl, or heteroarylalkyl, where the ring portion of each is

optionally substituted with one, two, three or four groups independently

selected from halogen, trifluoromethyl, trifluoromethoxy, cyano, nitro,

carboxyl, alkoxycarboxy, alkylcarboxy, hydroxy, lower alkyl, lower

alkoxy, amino, or mono or dialkylamino where each alkyl portion is lower

alkyl; or

R<sub>5</sub> and R<sub>6</sub> together with the carbon atom to which they are attached form a 5, 6, or 7

membered carbocyclic ring up to two of which members are optionally hetero

atoms selected from oxygen, sulfur and nitrogen.

Allowance of the claims and passage of the case to issue are respectfully solicited.

The Applicants urge the Examiner to contact the Applicants' undersigned representative at (312) 913-0001 if the Examiner believes that this would expedite prosecution of this application.

Respectfully submitted,

By: Stephen H. Docter  
Stephen H. Docter  
Reg. No. 44,659

Dated: January 3, 2005

McDonnell Boehnen  
Hulbert & Berghoff LLP  
300 South Wacker Drive  
Chicago, Illinois 60606  
(312) 913-0001